clinical transplant coordinator's responsibilities must include, but are not limited to, the following:

- (1) Ensuring the coordination of the clinical aspects of transplant patient and living donor care; and
- (2) Acting as a liaison between a kidney transplant center and dialysis facilities, as applicable.
- (d) Standard: Independent living donor advocate or living donor advocate team. The transplant center that performs living donor transplantation must identify either an independent living donor advocate or an independent living donor advocate team to ensure protection of the rights of living donors and prospective living donors.
- (1) The living donor advocate or living donor advocate team must not be involved in transplantation activities on a routine basis.
- (2) The independent living donor advocate or living donor advocate team must demonstrate:
- (i) Knowledge of living organ donation, transplantation, medical ethics, and informed consent; and
- (ii) Understanding of the potential impact of family and other external pressures on the prospective living donor's decision whether to donate and the ability to discuss these issues with the donor.
- (3) The independent living donor advocate or living donor advocate team is responsible for:
- (i) Representing and advising the donor;
- (ii) Protecting and promoting the interests of the donor; and
- (iii) Respecting the donor's decision and ensuring that the donor's decision is informed and free from coercion.
- (e) Standard: Transplant team. The transplant center must identify a multidisciplinary transplant team and describe the responsibilities of each member of the team. The team must be composed of individuals with the appropriate qualifications, training, and experience in the relevant areas of medicine, nursing, nutrition, social services, transplant coordination, and pharmacology.
- (f) Standard: Resource commitment. The transplant center must demonstrate availability of expertise in internal medicine, surgery, anesthesi-

ology, immunology, infectious disease control, pathology, radiology, blood banking, and patient education as related to the provision of transplantation services.

§ 482.100 Condition of participation: Organ procurement.

The transplant center must ensure that the hospital in which it operates has a written agreement for the receipt of organs with an OPO designated by the Secretary that identifies specific responsibilities for the hospital and for the OPO with respect to organ recovery and organ allocation.

§482.102 Condition of participation: Patient and living donor rights.

In addition to meeting the condition of participation "Patients rights" requirements at §482.13, the transplant center must protect and promote each transplant patient's and living donor's rights.

- (a) Standard: Informed consent for transplant patients. Transplant centers must implement written transplant patient informed consent policies that inform each patient of:
 - (1) The evaluation process;
 - (2) The surgical procedure;
 - (3) Alternative treatments;
- (4) Potential medical or psychosocial risks;
- (5) National and transplant center-specific outcomes, from the most recent SRTR center-specific report, including (but not limited to) the transplant center's observed and expected 1-year patient and graft survival, national 1-year patient and graft survival, and notification about all Medicare outcome requirements not being met by the transplant center;
- (6) Organ donor risk factors that could affect the success of the graft or the health of the patient, including, but not limited to, the donor's history, condition or age of the organs used, or the patient's potential risk of contracting the human immunodeficiency virus and other infectious diseases if the disease cannot be detected in an infected donor:
- (7) His or her right to refuse transplantation; and